UNITED STATES DISTRICT COURT DISTRICT OF MASSACHUSETTS

VERMONT PURE HOLDINGS, LTD.,)	
Plaintiff,)	
)	CIVIL ACTION NO
v.)	03-11465-DPW
)	
NESTLÉ WATERS NORTH)	
AMERICA, INC.,)	
Defendant.)	

MEMORANDUM AND ORDER March 28, 2006

Vermont Pure Holdings, Ltd. ("Vermont Pure") alleged in its original complaint that Nestle Waters North America, Inc. ("Nestle") violated §43(a) of the Lanham Act, 15 U.S.C. §1125(a)(1)(B), and multiple state law unfair competition statutes by making false or misleading statements in its advertising regarding the source, nature, and purity of Nestle's Poland Spring brand bottled water. In a Memorandum and Order dated September 9, 2004 ("September 9, 2004 Order"), 2004 WL 2030254 (D. Mass), I dismissed Vermont Pure's claims that Poland Spring water was not "spring water" on the grounds such claims were preempted by the Federal Food Drug & Cosmetic Act ("FDCA"), 21 U.S.C. §301, et seq. Id. at *7. The September 9, 2004 Order also denied Nestle's motion to dismiss with respect to the claims based on the origin and purity of the water ("quality claims"). Id. at *8.

Upon reference from me, Magistrate Judge Sorokin, who has been overseeing discovery in this case, issued Reports and

Recommendations ("R&R") with respect to the three motions now before me: (1) a motion by Vermont Pure for reconsideration of the September 9, 2004 Order in light of the Supreme Court's decision in Bates v. Dow Agrosciences, LLC, 125 S.Ct. 1788 (2005); (2) a motion by Nestle for the dismissal of the quality claims as presented in the Amended Complaint; and (3) a motion by Nestle to strike Vermont Pure's claim for disgorgement of profits, or in the alternative, enter judgment as a matter of law denying this form of relief.

Magistrate Judge Sorokin recommended that all three motions be denied. For the reasons discussed below, I accept this recommendation with respect to the latter two, but I find that my September 9, 2004 Order merits reconsideration and reversal of my earlier interlocutory order.

I. BACKGROUND

Drawing from Vermont Pure's original Complaint, the factual allegations of this case may be summarized as follows:

In 1854, Hiram Ricker began bottling spring water from a bedrock spring on Ricker Hill in Poland Maine. In the 1860's, the Ricker family opened a resort on the property, and in the 1880's, built a water bottling plant at the spring. The resort and bottled water sales thrived until the 1960s. By 1967, groundwater had ceased to flow out of the spring, and the bottling plant was closed.

In 1973, the Ricker family sold a 400-acre parcel of land surrounding the spring to a company called Waters of Maine.

Waters of Maine installed wells on its property and began to draw ground water from the Lower Range Pond Aquifer, which it marketed and sold as "spring water." Waters of Maine then built a new bottling plant on the property.

Perrier Co. ("Perrier") purchased the assets of Waters of Maine in 1979, and expanded the bottling plant. In a 1993 hostile takeover, Nestle S.A. purchased Perrier, and in 1994, further expanded the plant. Nestle also purchased Garden Spring Water Company, which was located several miles from the former Perrier plant. Nestle began bottling and marketing what it called Poland Spring water in approximately 1994.

Vermont Pure alleges that Poland Spring water has never been extracted from the Poland Spring, and does not even come from the same acquifer as the original source. Vermont Pure claims that Poland Spring water is pumped from the ground from a series of gravel-packed wells, some of which are located several miles from the original Poland Spring. Vermont Pure further alleges that Poland Spring derives this ground or well water not from "some of the most pristine and protected sources deep in the woods of Maine," as advertised, but from four publicly disclosed sources, and on occasion, from water trucked in from undisclosed out-of-state sources.

With respect to the purity of Poland Spring water, Vermont

Pure alleges that Nestle's production well techniques and methods

for withdrawing groundwater cause actual or potential

contamination of ground and well water.

II. DISCUSSION

28 U.S.C. §636(b)(1)(B) authorizes the parties to make timely objections to a magistrate's report and recommendation. This court then must make a <u>de novo</u> determination of those portions of the report to which objections are made. §636(b)(1)(B).

A. "Spring water" claims

Vermont Pure initially brought suit under §43(a) of the Lanham Act, 15 U.S.C. §1125(a)(1)(B), and multiple state unfair competition and false advertising statutes, alleging that Nestle's advertising for Poland Spring water was false or misleading because Poland Spring water was not "'spring water' in any regulatory, hydrological or plain meaning sense." Original Complaint at ¶15.

In the September 9, 2004 Order, I held that these claims were preempted by the FDCA. <u>Vermont Pure</u>, 2004 WL 3030254 at *7. In that decision, I noted that FDA regulations explicitly define the term "spring water." 21 C.F.R. §165.110(a)(2)(vi). While the FDCA clearly does not confer a private right of action to enforce its regulations directly, <u>see Mendes v. Medtronic, Inc.</u>, 18 F.3d 13, 19 n.4 (1st Cir. 1994), mere regulation of a term does not necessarily bar all false advertising claims relating to that term, <u>see Summit Tech.</u>, Inc. v. High-Line Med. Instruments Co., 933 F.Supp. 918, 933 (C.D. Cal. 1996). 2004 WL 3030254 at *3-*4.

I interpreted 21 U.S.C. §343-1's requirement of identity

between state and federal standards to mean that FDA regulations and requirements are exclusive and cannot be overriden by the states. 2004 WL 3030254 at *7.¹ I held that where, as here, the FDA has undertaken to define a term, the FDA has sole authority in misbranding cases relying upon that term. Drawing from Sandoz Pharmaceuticals v. Richardson-Vicks, 902 F.2d 222 (3rd Cir. 1990) and Braintree Labs, Inc. v. Nephro-Tech, Inc., 1997 WL 94237 (D. Kan. Feb. 26, 1997), I reasoned that whether a claim is preempted turns on whether it requires direct interpretation and application of the FDCA or FDA regulations. 2004 WL 3030254 at *4. Because Vermont Pure's claims required interpreting and applying the FDA's definition of "spring water", they were preempted. Id. at *7.

Vermont Pure seeks reconsideration only of the dismissal of its state law claims in light of the Supreme Court's April 2005 decision in <u>Bates</u>. A federal court may reconsider interlocutory orders and revise or amend them any time prior to final judgment. Fed R. Civ. P. 54(b); <u>Perez-Ruiz v. Crespo-Guillen</u>, 25 F.3d 40, 42 (1st Cir. 1994). In order to balance the competing interests of finality and justice, a motion for reconsideration should only

¹ 21 U.S.C. §343-1 provides, in pertinent part:

[[]N]o State or political subdivision of a state may directly or indirectly establish under any authority or continue in effect as to any food in interstate commerce --

⁽¹⁾ any requirement for a food which is the subject of a standard of identity established under section 341 of this title that is not identical to such standard of identity or that is not identical to the requirement of section 343(g) of this title...

²¹ U.S.C. §343-1(a)(1).

be granted if the movant demonstrates "(1) an intervening change in the law; (2) the discovery of new evidence not previously available; or (3) a clear error of law in the first order."

Davis v. Lehane, 89 F. Supp. 2d 142, 147 (D. Mass. 2000).

Vermont Pure contends that Bates presents a development in the relevant law and shows that the September 9, 2004 Order was clear error.

In <u>Bates</u>, Texas peanut farmers alleged that their crops were severely damaged by the application of Dow's "Strongarm" pesticide, which was registered by the Environmental Protection Agency (EPA) pursuant to its authority under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), 7 U.S.C. §136 et seq. <u>Bates</u>, 125 S.Ct. at 1790. The farmers threatened to sue Dow, alleging that Strongarm's label recommended its use in all peanut-growing areas when Dow knew or should have known that Strongarm would damage crops in soils with pH levels higher than 7.0. <u>Id</u>. In response, Dow sought a declaratory judgment that FIFRA preempted the farmers' claims. <u>Id</u>. The farmers then counterclaimed, bringing state law claims based on strict liability, negligence, fraud, and breach of warranty. <u>Id</u>. at 1790-91.

The Supreme Court granted certiorari to address the proper test for preemption in FIFRA cases. The Court began with a lengthy review of the legislative history of FIFRA, from which it concluded that "States have ample authority to review pesticide labels to ensure that they comply with both federal and state

labeling requirements." <u>Id.</u> at 1797. The Court then considered the text of 7 U.S.C. §136v, the provision at issue, which reads in relevant part:

(a) In general

A State may regulate the sale or use of any federally regulated pesticide or device in the State, but only if and to the extent the regulation does not permit any sale or use prohibited by this subchapter.

(b) Uniformity

Such State shall not impose or continue in effect any requirements for labeling or packaging in addition to or different from those required under this subchapter.

7 U.S.C. §136v. After carefully parsing subsection (b), the Court adopted a "parallel requirements" reading, which allows states to make and enforce labeling requirements so long as they are "equivalent to, and fully consistent with FIFRA's misbranding provisions." Id. at 1800. Thus, the Court held that "a state cause of action that seeks to enforce a federal requirement 'does not impose a requirement that is 'different from, or in addition to,' requirements under federal law.'" Id. (quoting Medtronic, Inc. v. Lohr, 518 U.S. 470, 513 (1996)).

The Court found unpersuasive Dow's arguments that the parallel requirements reading would lead to a "crazy-quilt" of anti-misbranding requirements put forth by juries in 50 states.

Id. at 1801. The Court held that FIFRA authorizes a relatively decentralized scheme, and private remedies enforcing FIFRA would further, rather than hinder, the functioning of the statute. Id. at 1802.

Magistrate Judge Sorokin rejected Vermont Pure's request that this court follow <u>Bates</u> and adopt a parallel requirements

reading of 21 U.S.C. §343-1. His decision focused on the differences between the nature of the regulatory schemes established by FIFRA and the FDCA. He noted that even before Bates, the Supreme Court had found that FIFRA did not occupy the field to the exclusion of the states. September 23, 2005 R&R at 3 (citing Wisconsin Public Intervenor v. Mortier, 501 U.S. 597, 607 (1991)). In his view, Bates merely confirmed the narrow preemptive effect of FIFRA by holding that §136v allows the imposition of state sanctions for violating state rules that duplicate federal standards. September 23, 2005 R&R at 4; Bates, 125 S.Ct. at 1797.

In contrast, Magistrate Judge Sorokin found that the FDA mandates that state requirements be barred unless they are "identical" to the federal requirements, 21 U.S.C. §343-1, and that all enforcement be centralized in the FDA, 21 U.S.C. § 337(a). September 23, 2005 R&R at 4. Section §337, which has no equivalent in FIFRA, requires that all enforcement actions be brought by the federal government and has only one narrow exception allowing states to bring actions in their own names. 21 U.S.C. § 337(a), (b). From these provisions, Magistrate Judge Sorokin concluded that "Congress established the FDA as the interpreter and enforcer of the FDCA." September 23, 2005 R&R at 5.

Magistrate Judge Sorokin acknowledged that §343-1 permits states, in at least some circumstances, to impose requirements "identical" to federal requirements and to enforce or create

remedies for those regulations. <u>Id.</u> However, based on what he viewed as a marked difference between the role of the EPA in enforcing FIFRA and the FDA in enforcing the FDCA, he rejected the notion that <u>Bates</u> amounted to "an intervening change in controlling law" or "evidence of clear error of law" sufficient to prompt a reconsideration of the September 9, 2004 Order. <u>Id.</u>

Vermont Pure objects to Magistrate Judge Sorokin's decision on two grounds. First, it claims that he misunderstood the text of §343-1. Vermont Pure argues that FIFRA's §137v is substantively identical to §343-1 in that they each allow states to impose requirements in federally regulated areas, so long as those requirements are identical to the federal ones. Docket No. 122 at 8. Invoking the long-settled principle that courts may only find preemption when congressional intent is "clear and manifest", <u>Bates</u>, 125 S. Ct. at 1801, Vermont Pure argues that the text of §343-1, like that of FIFRA in <u>Bates</u>, expressed no such preemptive intent, either explicitly or implicitly.

Second, Vermont Pure contends that Magistrate Judge Sorokin misconstrued the scope of the FDCA's regulatory scheme. It claims that the FDCA has never been held to occupy the field of food regulation, and as evidenced by the FDA's web site, the FDA relies on state regulation in the area of bottled water. Docket No. 122 at 11-12 (citing http://www.cfsan.fda.gov/~dms/botwatr.html). Vermont Pure points to the legislative history of § 343-1, which it claims suggests a limited preemptive effect.

After carefully considering the Court's ruling in Bates, I

find that I must reconsider my holding in the September 9, 2004

Order and the test that I employed for preemption under the FDCA.

I begin with the relevant text of §343-1:

- (a) [N]o State or political subdivision of a state may directly or indirectly establish under any authority or continue in effect as to any food in interstate commerce --
- (1) any requirement for a food which is the subject of a standard of identity established under section 341 of this title that is not identical to such standard of identity or that is not identical to the requirement of section 343(g) of this title...

21 U.S.C. §343-1(a)(1). The introductory words "[N]o State or political subdivision of a state" clearly establish the intent of Congress to entirely occupy at least a portion of the field of food regulation. The more difficult task is to describe what portion and delineate the scope of the preemptive effect expressed in Paragraph (a)(1).

The term "requirement" "sweeps broadly." <u>Cipollone v.</u>

<u>Ligget Group, Inc.</u>, 505 U.S. 504, 521. Yet, it is not so

expansive as to include "[a]n occurrence that merely motivates an optional decision," such as a jury verdict that might induce a pesticide maker to change its label; rather it incorporates

"positive enactments, such as statutes and regulations, as well as common-law duties." <u>Bates</u>, 125 S.Ct. at 1798.

Section 343-1(a)(1) relates only to those state requirements for foods that are the "subject of a standard of identity established under section 341 of this title." 21 U.S.C. §343-1(a)(1). A "standard of identity" is a regulation in which the FDA fixes the ingredients of, and consequently the definition of,

a food. 62 Cases, More or Less, Each Containing Six Jars of Jam v. U.S., 340 U.S. 593, 598 (1951). There is no doubt that the FDA has promulgated such a regulation with respect to bottled water in general, and spring water in particular. See 60 Fed. Reg. 57076, 57076 (Nov. 13, 1995); 21 C.F.R. §165.110.

Section 341(a)(1) states that "no State" may establish requirements that are "not identical to such standard of identity" or that are "not identical to the requirement of section 343(g)."² 21 U.S.C. §343-1(a)(1). The negative pregnant evident from a plain reading of this language is that states may establish requirements that are identical to FDA standards. The term "identical," does not refer to the specific words used in the state requirement, but to the direct and indirect effects of the requirement. See 21 C.F.R. 100.1(c)4. In this connection, I find and adopt as my own a definition of "identical" which has been incorporated in pending legislation to modify the FDCA: "the term 'identical' means that the language is substantially the same language as the comparable provision of the [FDCA], and that any difference does not result in the imposition of materially

² Section 343(g) provides that:

A food shall be deemed to be misbranded... [i]f it purports to be or is represented as a food for which a definition and standard of identity has been prescribed by regulations as provided by section 341 of this title, unless (1) it conforms to such definition and standard, and (2) its label bears the name of the food specified in the definition and standard, and, insofar as may be required by such regulations, the common names of optional ingredients (other than spices, flavoring, and coloring) present in such food.

²¹ U.S.C. §343(q).

different requirements." H. Rep. No. 109-379 (Feb. 28, 2006).

Applied in the context of this case, the plain meaning of §343-1(a)(1) is that states may regulate bottled water so long as their standards mirror those set by the FDA. Thus, the analysis for express preemption turns not on whether a claim requires direct interpretation and application of FDA regulations, as I stated in the September 9, 2004 Order, but on the identity of the state and federal regulations. See Bates, 125 S.Ct. at 1800.

FDA regulations define "spring water" as "water derived from an underground formation from which water flows naturally to the surface of the earth." 21 C.F.R. §165.1110(a)(2)(vi). To the extent the state laws under which Vermont Pure brings suit have adopted, by recitation or reference, the FDA's definition of "spring water," they are not expressly preempted by §343-1.

³ Section 337 does not change this analysis. Section 337(a) clearly states that only the federal government can bring an action to enforce the FDCA. 21 U.S.C. §337(a). Section 337(b) creates an exception for states to enforce certain provisions, including §343(g)'s standard of identity requirements. 21 U.S.C. §337(b). These provisions, however, relate only to enforcement of the FDCA. Vermont Pure's "spring water" claims arise purely from state law. To be sure, the state laws are based on federal regulations, as the FDCA requires them to be, but Section 337 is silent as to these state law suits. This reading makes sense not only in terms of the plain language of §337, but in light of the legislative history and the agency's interpretive history of §343-1, see infra.

⁴ FDA requirements also specify in detail the method of extraction to be used for acquiring spring water. 21 C.F.R. §165.110(a)(2)(vi).

 $^{^{5}}$ In Count II of its original complaint, Vermont Pure alleged violation of the unfair competition statutes of least twenty-five states. In its objections to Magistrate Judge Sorokin's R&R, it lists only six states. Docket No. 122 at 10, n.5. Each of these states appears to have incorporated the FDA

Although the text of §343-1(a)(1) is strong evidence that Congress did not expressly preempt Vermont Pure's state law claims, Nestle nevertheless contends that they are impliedly preempted. An examination of the legislative history and the agency's interpretive history lead to the opposite conclusion.

When Congress passed §343-1 as part of the Nutrition
Labeling and Education Act of 1990 (NLEA), it expressly
considered the question of preemption. The purpose of the NLEA
was not to preclude all state regulation of nutrition labeling,
but to "prevent State and local governments from adopting
inconsistent requirements with respect to the labeling of
nutrients." H. Rep. No. 101-538, 1990 U.S.C.C.A.N. 3336,
3337 (emphasis added). In service of this purpose, Congress

definition of "spring water." <u>See</u> Conn. Gen. Stat. Ann. §21a-150 ("natural water obtained from an underground formation from which water flows naturally to the surface of the earth"); 105 Mass. Regs. Code §570.012(C)(2)(a) (explicitly adopting definition in 21 C.F.R. §165.1110(a)(2)(vi)); Code Me. R. 10-144 Ch. 235, §2 ("Water derived from an underground formation from which water flows naturally to the surface of the earth"); N.H. Code Admin. R. Env-Ws 389.03, HE-P 2101.01 (explicitly adopting definition in 21 C.F.R. §165.1110(a)(2)(vi)); N.Y. Comp. Codes R. & Regs. tit. 10, §5-6.3 ("water derived from an underground formation from which water flows naturally to the surface of the earth"); R.I. Code R. 14 180 007 ("water derived from an underground formation from which water flows naturally to the surface of the earth").

Vermont Pure has not provided an amended complaint describing under which state laws it intends to proceed. Nestle argues that Vermont Pure should not be granted leave to amend given the status of discovery and the extensive defense preparation that has taken place thus far. Docket No. 131 at 19, n.11. Fed. R. Civ. P. 15(a) is a "liberal amendment policy" and because I find no "undue delay" or "undue prejudice," Invest Almaz v. Temple-Inland Forest Products Corp., 243 F.3d 57, 71(1st Cir. 2001), I will permit Vermont Pure to file a Second Amended Complaint identifying in Count II precisely the state law upon which it will seek to establish a "spring water" claim.

expressly declared "[t]he Nutrition Labeling and Education Act of 1990 shall not be construed to preempt any provision of State law, unless such provision is expressly preempted under section 403A of the [FDCA]." Pub. L. 101-535, 104 Stat 2353, 2364 (Nov. 8, 1990).

The FDA echoed this view when it promulgated its final rule interpreting §343-1. In response to requests for clarification as to preemptive scope, the FDA stated "if the State requirement is identical to the Federal law, there is no issue of preemption... [T]he only State requirements that are subject to preemption are those that are affirmatively different from the Federal requirements." 60 C.F.R. 57076-01, 57120 (Nov. 13, 1995). The FDA also clarified that where no federal requirement exists, preemption does not occur. Id.

Nestle argues that the FDCA creates a "thorough, comprehensive and precise regulatory scheme" and the goal of uniform national standards mandates preemption of Vermont Pure's claims. It points to the technical extraction criteria for "spring water" listed in 21 C.F.R. 165.110(a)(2)(vi) and Congress' stated goal of relieving industry from "some types of State requirements that interfere with [industry's] ability to market products in all 50 States in an efficient and costeffective manner." 60 C.F.R. 57076-01, 57120 (Nov. 13, 1995). The specificity and uniformity of the definition of "spring"

⁶ Section 403A of the Nutrition Labeling and Education Act of 1990 is codified as 21 U.S.C. §343-1.

water," however, are not directly threatened by a claim under state deceptive practices law where, as here, it appears that at least some of the state laws forming the basis for the lawsuit have adopted the federal definition of "spring water."

This case is also distinct from a situation in which a state enacts a statute that substantively adds to federal labeling requirements. As evidence of the importance of uniformity, Nestle points to the FDA's opinion that a Maine statute that required labels on bottled water to identify the name and geographic location of the source of the water was preempted. Here, Vermont Pure does not seek either to challenge or add to the FDA's definition of "spring water."

Nestle contends that Vermont Pure's claims have the indirect effect of adding to the FDA regulations by imposing the "requirement" that Nestle prove in private litigation that its bottled water meets the "spring water" standard set by the FDA. Nestle also argues that the claims add the requirement of various state statutory remedies, such as enjoining labeling and advertising, damages, and attorney's fees. Justice Thomas touched on a similar concern in his opinion concurring in part and dissenting in part in Bates: "[a] state-law cause of action, even if not specific to labeling, nevertheless imposes a labeling requirement 'in addition to or different from' FIFRA's when it attaches liability to statements on the label that do not produce liability under FIFRA." Bates, 125 S.Ct. at 1805. However, the language of §343-1 is clear that state remedies for failing to

meet FDA standards of identity are not "requirements" and are not foreclosed.

I recognize that allowing Vermont Pure's state law claims arguably threatens the goal of uniform interpretation. There is the prospect that judges and juries in 50 different states interpreting a precise FDA definition and then imposing injunctions and damages on a bottled water producer will generate inconsistencies in interpretation of the identical provision.

But Congress and the FDA appear to have made a conscious choice to allow the several states to regulate bottled water so long as the state standards employed are identical to those adopted by the FDA. The requirement of identity promotes uniformity in that courts in every state look to the same standard. Permitting states to interpret FDA definitions also allows for additional resources to be brought to bear in the enforcement of national standards upon regional and local markets.

Having been instructed by the Supreme Court's approach in Bates, I cannot ignore the text, legislative history, and the agency's interpretive guidance regarding §343-1. I find that Vermont Pure's "spring water" claims are not preempted to the extent that they are based solely on state laws that provide private rights of action and have adopted the FDA definition of

⁷ Of course, I rely on <u>Bates</u> only by analogy for my adoption of a parallel requirements reading of §343-1. I take from <u>Bates</u> the general model for statutory interpretation in the context of legislation implicating a theory of parallel federal requirements, but my conclusions as to the issue at hand derive from consideration of the FDCA itself.

"spring water."

B. Quality claims

Vermont Pure's Amended Complaint alleges a single count of false advertising under §43(a) of the Lanham Act. Amended Complaint at ¶¶ 38-46. Vermont Pure contends that Nestle falsely or misleadingly markets Poland Spring water as "pure, even though Poland Spring is aware of contamination, and the potential for contamination of its water sources." Id. at ¶ 13f. In this connection, Vermont Pure claims that test samples of Poland Spring water have revealed excessive levels of Heterotrophic Plane Count ("HPC") bacteria, excessive chlorination, and contamination by fecal coliform. Id. at ¶¶ 21, 24. Plaintiff also claims that Nestle's extraction practices "have the actual or potential ability to extract contaminated or potentially contaminated water." Id. at ¶ 26.

The Amended Complaint further alleges that in order to compete with Vermont Pure, Nestle devised a strategy to market its water as "pure water from a natural protected source in Maine," even though it was neither pure nor from a natural source in Maine. Id. at ¶ 35. Vermont Pure asserts that, consistent with this strategy, Nestle made the following false or misleading statements in its commercial advertising: that Poland Spring bottled water is "filtered and purified deep underground;" that it comes from "some of the most pristine and protected sources deep in the woods of Maine;" it "remains unspoiled because thousands of guards protect it;" it "only comes from one source -

- in the wilderness of Maine;" and it "comes straight from nature, straight from Maine." <u>Id.</u> at \P 41. Vermont Pure claims that the name "Poland Spring" is misleading because it suggests that the water actually comes from the Poland Spring, and in 2002, Nestle changed its label to read "Poland Spring Brand" in recognition of the "material misrepresentation in its advertising." <u>Id.</u> at \P 41(e), (g).

Pursuant to Fed. R. Civ. P. 12(c), Nestle has moved to dismiss the claims in Plaintiff's Amended Complaint relating to the quality, purity, treatment, testing, and contamination of Poland Spring's water sources. Docket No. 98. Magistrate Judge Sorokin recommended that I deny the motion. December 12, 2005 R&R. I will adopt his recommendation.

In reviewing a motion for judgment on the pleadings under Rule 12(c), "the district court must accept all of the nonmoving party's well-pleaded factual averments as true and draw all reasonable inferences in her favor." Feliciano v. Rhode Island, 160 F.3d 780, 788 (1st Cir. 1988). The motion may not be granted "unless it appears beyond a doubt that the nonmoving party can prove no set of facts in support of her claim which would entitle her to relief." Id.

Magistrate Judge Sorokin rejected Nestle's argument that the quality claims were preempted by the FDCA. First, he found that unlike Plaintiff's claims regarding "spring water", the quality claims do not arise out of a term that is specifically defined by the FDA. December 12, 2005 R&R at 5. "Indeed," he

explained, "the FDA expressly declined to 'use its resources to define the term 'pure,'' a term central to Plaintiff's amended claims." <u>Id.</u> (quoting 60 Fed. Reg. 57076, 57099 (1995)).

Magistrate Judge Sorokin's holding turned on the finding that Vermont Pure's claims question the truth of Nestle's advertising, not whether Poland Spring water meets FDA standards.

Id. He rejected the argument that the FDCA so comprehensively regulates the field that a plaintiff may never bring a Lanham Act claim regarding the misleading advertising of bottled water quality, purity, testing, treatment, and sources. Id. at 6.

Magistrate Judge Sorokin distinguished the cases upon which Nestle relied for its preemption argument. He focused on <u>Sandoz Pharmaceuticals v. Richardson-Vicks</u>, 902 F.2d 222 (3rd Cir. 1990), which Nestle cited in support of a broad view of the FDCA's preemptive effect. <u>Sandoz</u> involved a claim of false advertising based upon the use of the term "inactive" with respect to demulcents. <u>Id.</u> at 230. The <u>Sandoz</u> court refused to usurp what it viewed as the FDA's responsibility to define the term "inactive" and decide whether the demulcent at issue fit that definition. <u>Id.</u> at 231. Magistrate Judge Sorokin found that Vermont Pure's claims, in contrast to those in <u>Sandoz</u>, require no interpretation, application, or enforcement of FDA regulations.

Similarly, he found inapplicable <u>Mylan Laboratories</u>, <u>Inc. v.</u>

<u>Matkari</u>, 7 F.3d 1130, 1139 (4th Cir. 1993) (dismissing as preempted by the FDCA a claim that merely placing a drug on the

market is a statement of false FDA approval actionable under the Lanham Act), and <u>Animal Legal Defense Fund v. Promivi Veal Corp.</u>, 626 F. Supp. 278, 283-84 (D. Mass. 1986) (finding that the FDCA and the Federal Meat Inspection Act preempted state law actions forcing veal producers to label their products as "adulterated" and to add to the label bacteria content, use of antibiotics, and genetic alteration information).8

Nestle objects to Magistrate Judge Sorokin's R&R arguing that he erred in his interpretation of the scope of federal preemption. First, Nestle contends that Magistrate Judge Sorokin mistakenly relied on Vermont Pure's statement in its brief that Nestle made false claims in order to differentiate Poland Spring from other bottled water. Docket No. 155 at 8. Nestle argues that the heart of Vermont Pure's claims is not statements in Nestle's advertising comparing its product to its competitors' products, but the quality of Poland Spring water as measured by a variety of standards. Id. at 8,10. Because the FDA sets forth what Nestle labels "exclusive quality standards," evaluation of water purity by any other standard, according to Nestle, encroaches upon the FDA's authority. Id. Similarly, Nestle claims that Vermont Pure's intention to consult FDA and non-FDA

⁸ Because Nestle advanced different arguments in the instant motion to dismiss than those pressed in the prior motion to dismiss (Docket No. 32), Magistrate Judge Sorokin recommended against simply applying the Law of the Case doctrine to bar the present motion. December 12, 2005 R&R at 5 n.4. I agree that the prior motion raised arguments that are distinct from those under consideration here, and I decline to deny the motion on the basis of the Law of the Case doctrine.

standards for allowable contaminate levels also flouts the federal regulatory scheme. ⁹ Id. at 11-12.

Nestle further argues that Magistrate Judge Sorokin's recognition that the FDA has authority over the term "pure" leads to the conclusion that the FDA, and not the courts, should define that term. Nestle points to two courts that deferred to the FDA on what it contends are analogous questions: Sandoz, 902 F.2d at 231 (leaving to the FDA the question of whether a demulcent should be labeled "active" or "inactive"), and Abruzzi Foods, Inc. v. Pasta & Cheese, Inc., 687 F. Supp. 20, 21 (D. Mass. 1988) (holding that "this court lacks the authority to create a standard of what constitutes "freshness" in the pre-packaged

⁹ Nestle has submitted to the court Vermont Pure's answers to interrogatories stating the standards that Vermont Pure intends to use to prove its case. Magistrate Judge Sorokin did not consider the interrogatory answers in his decision because, although they were available at the time the R&R was issued, they were not provided when the parties briefed the motion. December 12, 2005 R&R at 9, n.9. Nestle argues that not only should Magistrate Judge Sorokin have considered the interrogatories, but because they constitute "matters outside the pleadings," the motion to dismiss should have been converted to a motion for summary judgment. Docket No. 155 at 11; Fed. R. Civ. P. 12(c).

Rule 12(c) gives a court the option of considering the evidence and converting to summary judgment, or excluding it and deciding the motion for judgment on the pleadings. Fed. R. Civ. P. 12(c). See Garita Hotel Ltd. Partnership, Etc. v. Ponce Federal Bank, F.S.B., 958 F.2d 15, 18 (1st Cir. 1992) ("If the district court chooses to ignore the supplementary materials and determines the motion under the Rule 12(b)(6) standard, no conversion occurs."). Magistrate Judge Sorokin did not consider the interrogatories and properly treated the motion as a judgment on the pleadings. I do the same here. At this stage in the litigation, with discovery still ongoing, I will not convert a Rule 12(c) motion in a complex case based on the belated submission of a single set of interrogatory answers.

pasta industry"). 10

In short, Nestle argues that the quality claims cannot be proven without interpretation and application of FDCA standards, and therefore, they are preempted. I disagree. No federal standards of identity for bottled water purity exist. It is clearly within the FDA's power to promulgate such standards, but as Magistrate Judge Sorokin noted, it has chosen not to do so. 60 Fed. Req. 57076 at 57099.

Nestle asserts that the failure of the FDA to regulate a term precludes Lanham Act actions based upon that term. To the contrary, as the FDA observed, "[i]f there is no Federal requirement to be given preemptive effect, preemption does not occur." Id. at 57120. The fact that the FDA sets standards for contaminates in bottled water, including the chlorine and bacteria that Vermont Pure alleges have been found in Poland

Nestle also argues that Magistrate Judge Sorokin raised sua sponte a possible distinction in the FDCA's misbranding regulations between advertising and labeling, and the parties never had the opportunity to brief the issue. Docket No. 155 at 15. Vermont Pure responds that Nestle failed to raise this point with the Magistrate and is barred from raising it now. Docket No. 161 at 14. Because the parties did not raise or brief this issue before Magistrate Judge Sorokin and because I decide on other grounds whether Vermont Pure's claims are preempted, there is no need to delve into the question of whether the FDCA's misbranding regulations extend to advertising as well as labeling.

If I were to consider that issue, I would observe that 21 U.S.C. §§334(a) and 321 clearly include advertising as within the scope of the FDCA's definition of "misbranded." However, these provisions do not apply to the issue in this case. The question here is not whether Poland Spring water is misbranded for purposes of the FDCA, but whether Nestle has falsely or misleadingly represented the purity and source of Poland Spring water.

Spring water, does not change this conclusion. In the absence of a federal standard of identity, private litigants may look to other standards, including FDA standards, to interpret the term at issue.

The First Circuit's opinion in <u>Abruzzi</u>, a case discussed by both Vermont Pure and Nestle, is instructive. 986 F.2d 605 (1st Cir. 1993). In <u>Abruzzi</u>, a pasta producer and wholesaler sued a competitor for calling its pasta "fresh" allegedly in violation of Massachusetts' unfair competition laws. <u>Id.</u> at 605. The term "fresh" as applied to refrigerated pasta had not been defined in any federal or state law, statute, or regulation. The District Court granted summary judgment on the grounds that only the FDA or a legislative body has the authority to define the term. 687 F. Supp. at 21, n.2.

The First Circuit affirmed the District Court, but on entirely different grounds. Without reaching or even mentioning the question of a court's competence to define food-related terms, then-Chief Judge Breyer reached the merits of the case. He asked whether Abruzzi had set forth facts sufficient to show that its use of the term "fresh" was deceptive. 986 F.2d at 605. In answering that question, he looked to various sources for a definition of "fresh": FDA rules for milk, FDA rules on the use of the term "fresh", magazine articles, even "common sense." Id. at 606. With respect to the fact that the FDA specifically declined to define "fresh," Chief Judge Breyer held that Abruzzi must look elsewhere for support. Id. He did not find Abruzzi's

claim preempted merely because the FDA had the (unexercised) authority to define the term at issue.

Magistrate Judge Sorokin appropriately distinguished <u>Sandoz</u>, <u>Promivi</u>, and <u>Mylan Labs</u>, the three cases upon which Nestle principally relies. I add only that the FDA has in this setting specifically refused to define the term "pure." 60 Fed. Reg. at 57099. This is in contrast to <u>Sandoz</u>, 902 F.2d at 230-31, in which the plaintiff asked the court to decide if a defined FDA term ("active") applied to a particular product (a demulcent); <u>Promivi</u>, 626 F. Supp. at 283, in which the FDCA, in conjunction with the Federal Meat Inspection Act, was found to have created a comprehensive federal scheme controlling medicated animal feeds, drugs used to treat animals raised for human consumption, and meat labeling, packaging, and marketing; and <u>Mylan Labs</u>, 7 F.3d at 1139, in which the plaintiff alleged that the defendants falsely represented FDA approval of a product.

Although Magistrate Judge Sorokin did not address it in detail, Lynnbrook Farms v. Smithkline Beecham Corporation, 79

F.3d 620 (7th Cir. 1996), is also inapposite. In Lynnbrook Farms, a farm partnership sued a cattle vaccine manufacturer when the vaccines proved ineffective and harmful to cattle. The case raised the question whether the United States Department of Agricultures's (USDA) declaration of preemption, made through Animal and Plant Health Service (APHIS) pursuant to the federal Virus-Serum-Toxin Act (VSTA), 21 U.S.C. §§151-159, preempted the plaintiff's numerous state law claims. The preemption clause

promulgated by APHIS provided:

states are not free to impose requirements which are different from or in addition to, those imposed by USDA regarding the safety, efficacy, potency or purity of a product

57 Fed. Reg. 38759 (Aug. 27, 1992). The Seventh Circuit interpreted this provision as "just another way of saying a [s]tate shall not impose... any requirements." 79 F.3d at 628 (quoting Cipollone, 505 U.S. at 371). That interpretation was consistent with readings given by several other courts of nearly identical language in FIFRA and the Medical Device Amendments to the FDCA (MDA). Lynnbrook Farms, 79 F.3d at 628-69. It was also consistent with the intent of APHIS, which expressly stated that the regulation was designed to preempt state tort actions when the defendant complied with federal regulations, but to allow them when the defendant ignored or violated federal rules. Id. at 629.

Lynnbrook Farms differs significantly from the instant case in that APHIS clearly explained that it intended to preempt the state actions in question. Here, the FDA did the opposite: it expressly declined to regulate the term "pure." More fundamentally, the approach to interpretation of the preemption clause in Lynnbrook Farms is suspect after Bates, 125 S.Ct. at 1800, which adopted a parallel requirements reading of the "in addition to or different from" language in FIFRA.

Nothing in the text, legislative history, agency interpretation or caselaw of the FDCA suggests that it preempts Vermont Pure's claims regarding Nestle's allegedly misleading

representations regarding the quality, purity, treatment, contamination and source of Poland Spring water. Accordingly, I accept Magistrate Judge Sorokin's recommendation to deny Nestle's motion to dismiss.

C. Disgorgement of profits

In its Amended Complaint, Plaintiff includes the following prayer for relief: "all of Nestle's profits and/or pecuniary gains derived directly or indirectly from Nestle's unlawful conduct, as permitted by 15 U.S.C. §1117." Amended Complaint at 16. Nestle argues that the remedy of disgorgement of profits is not available to Plaintiff in this case as a matter of law.

Accordingly, it has moved to strike pursuant to Fed. R. Civ. P. 12(f), or in the alternative for judgment as a matter of law as allowed by Rule 12(c). Docket No. 74. Magistrate Judge Sorokin recommended that I deny Defendant's motion. Again, I accept this recommendation.

Magistrate Judge Sorokin found that contrary to Nestle's contention, the Lanham Act does not, as a matter of law, preclude the remedy of disgorgement of profits in a non-comparative false

¹¹ As a threshold matter, I note that Nestle has adequately established that its Objection to the September 20, 2005 R&R was timely raised. This issue not previously addressed by Nestle despite its inclusion at the outset of Vermont Pure's opposition to the Objection was raised by me at the hearing in this matter where Nestle was given the opportunity to respond with a post hearing submission. That submission establishes -- and Vermont Pure does not now dispute -- that despite the date of the September 20, 2005 R&R, the objection was served within ten days from service of the R&R on Nestle. See generally Fed. R. Civ. P. 72.

advertising case. September 20, 2005 R&R at 7. Looking first to the statutory text, he observed that 15 U.S.C. §1117(a), the provision governing damages for Lanham Act cases, specifically allows for plaintiffs to recover defendants' profits, subject to principles of equity and the discretion of the court. Id. at 4-5. He then noted that in Aktiebolaget Electrolux v. Armatron Int'l Inc., 999 F.2d 1, 5 (1st Cir. 1999), the First Circuit required a plaintiff seeking disgorgement to prove both actual harm and direct competition. In the absence of those elements, damages may be awarded upon one of three conditions: the defendant acted fraudulently; to avoid unjust enrichment of the defendant; or to deter further willful misconduct. Id. In Tamko Roofing Products v. Ideal Roofing Co., 282 F.3d 23, 36 (1st Cir. 2002), the First Circuit described three justifications for awarding an accounting:

(1) as a rough measure of the harm to plaintiff; (2) to avoid unjust enrichment of the defendant; of (3) if necessary to protect the plaintiff by deterring a willful

¹² Magistrate Judge Sorokin relied on the following passage in which the Court stated four rules for awarding monetary damages under §1117:

¹⁾ a plaintiff seeking damages must prove actual harm, such as the diversion of sales to the defendant; 2) a plaintiff seeking an accounting of defendant's profits must show that the defendant's profits would have gone to plaintiff if there was no violation; 3) the general rule of direct competition is loosened if the defendant acted fraudulently or palmed off inferior goods, such that actual harm is presumed; and 4) where defendant's inequitable conduct warrants bypassing the usual rule of actual harm, damages may be assessed on an unjust enrichment or deterrence theory.

<u>Aktiebolaget Electrolux v. Armatron Int'l Inc.</u>, 999 F.2d 1, 5 (1st Cir. 1999).

infringer from further infringement.

In its Amended Complaint, Vermont Pure alleged that Nestle's alleged violations proximately caused a direct diversion of sales from Vermont Pure to Nestle. Amended Complaint at \P 46. Noting that to satisfy Rule 12(c), Vermont Pure need only allege one set of facts for which disgorgement is available, Magistrate Judge Sorokin concluded that although far from making the factual showing necessary for a damage award, Vermont Pure's allegations of actual harm and direct competition were sufficient to overcome a judgment on the pleadings. September 20, 2005 R&R at 7.

Nestle objects to this decision on the grounds that Magistrate Judge Sorokin misconstrued the legal standard. Specifically, it takes issue with the following statement:

Vermont Pure might obtain Nestle's profits as one form of relief in this case if, after prevailing on the merits of the claim that Nestle engaged in false advertising under the Lanham Act, it establishes actual harm and direct competition <u>such that</u> Nestle's profits would have been Vermont Pure's.

R&R at 5 (emphasis in original).

Nestle argues that in a non-comparative advertising case, such as this one, the plaintiff must include a showing of "culpable conduct," such as intentional deception and targeted wrongdoing, on the part of the defendant in order to merit disgorgement of profits. Nestle's Objection at 2 (Oct. 5,

As Magistrate Judge Sorokin noted, there is no dispute that this is a non-comparative advertising case, as the challenged advertising makes no direct comparison between Poland

2005). Magistrate Judge Sorokin, it claims, based his recommendations on principles derived from cases involving trademark infringement, palming off, or other actions that directly compare defendant's and plaintiff's products. Since the First Circuit has never directly ruled on the availability of disgorgement of profits in a non-comparative advertising case, Nestle urges this court to follow Eckel Indus., Inc. v. Primary Bank, 26 F. Supp. 2d 313 (D.N.H. 1998) and ALPO Petfods, Inc. v. Ralston Purina Co., 913 F.2d 958 (D.C. Cir. 1990), among others, in holding that the remedy of disgorgement of profits in false advertising cases is limited to cases involving comparative advertising or deceptive acts that would lead consumers to mistake the defendant's product for the plaintiff's. See Eckel, See Eckel, Supp. at 318.

I decline the invitation to embrace a per se rule forbidding the remedy of disgorgement of profits in non-comparative advertising cases. I begin, as did Magistrate Judge Sorokin, with the text of the governing statute. Section 1117(a) provides that when a violation of §1125(a) has been established,

the plaintiff shall be entitled, subject to the provisions of sections 1111 and 1114 of this title, and subject to the principles of equity, to recover (1) defendant's profits, (2) any damages sustained by the plaintiff, and (3) the costs of the action.

Spring and any of its competitors. September 20, 2005 R&R at 4. Although Vermont Pure's allegations that Nestle conducted an "ongoing, pervasive fraud on American consumers", Amended Complaint at \P 12, suggests bad faith, Vermont Pure made no specific allegations that Nestle made false statements knowingly, willfully, or in bad faith. \underline{Id} .

15 U.S.C. §1117(a). This provision does not limit or suggest limiting damages on the basis of the comparative or non-comparative nature of the false advertising claim. The three types of allowable damages are restricted only by §1111, which requires notice to the defendant; §1114, which specifically limits remedies for copyright infringement; "principles of equity;" and the discretion of the court "according to the circumstances of the case." §1117(a).

Case law supports a case-by-case determination of available remedies. As Magistrate Judge Sorokin pointed out, the standard articulated by First Circuit in AB Electrolux and Tamko focuses on actual harm, direct competition, and equitable considerations; bad faith is not necessarily required. September 20, 2005 R&R at 7; AB Electrolux, 999 F.2d at 5; Tamko, 282 F.3d at 36. The Tamko court specifically distinguished the rule of the First Circuit from the Restatement on the question of bad faith: "it has been this circuit's rule that an accounting of defendant's profits where the products directly compete does not require fraud, bad faith, or palming off." Tamko, 282 F.3d at 36. Put simply, AB Electrolux and Tamko teach that actual harm and direct competition are sufficient to support a damages award in a Lanham

The <u>Tamko</u> court explicitly left unresolved the question of whether "willfulness" is a precondition for an accounting. <u>Tamko Roofing Products</u>, <u>Inc. v. Ideal Roofing Co.</u>, Ltd., 282 F.3d 23, 36 (1st Cir. 2002).

Act case. 15 Id. And because it is an equitable remedy, determination of such an award should be made on a case-by-case basis.

Nestle distinguishes <u>AB Electrolux</u> and <u>Tamko</u> on the grounds that they were trademark infringement cases, and consequently, their holdings do not apply to non-comparative false advertising claims. Nestle's Objection at 6-8. I disagree. In both cases, the First Circuit used generic language to articulate broad Lanham Act principles. Throughout the opinions, the Court makes general statements about the nature of the Lanham Act that apply beyond the infringement context. <u>See e.g., Tamko</u>, 282 F.3d at 38 ("Mechanical rules are of little aid in this analysis [of damages]"; "Equity must take account of the purposes served by the Lanham Act"). I decline to ignore this guidance, especially when the First Circuit has yet to address the specific question at hand.

Nestle relies heavily on <u>Eckel Indus., Inc. v. Primary Bank</u>, 26 F. Supp. 2d 313 (D.N.H. 1998), for the proposition that disgorgement of profits is not available for non-comparative false advertising claims. In <u>Eckel</u>, a manufacturer of impact traffic doors brought suit under the Lanham Act against a competitor for allegedly using photos of the plaintiff's doors in defendant's advertising materials and identifying them as

¹⁵ To be sure, actual harm and direct competition may be more difficult to prove in a non-comparative false advertising case than for example, in an infringement case.

defendant's. <u>Id.</u> at 315. Eckel alleged that the defendant made false and misleading statements about Eckel's products and "palmed off" its doors as Eckel's doors. <u>Id.</u> at 316.

In applying the law as stated in <u>AB Electrolux</u>, the <u>Eckel</u> court questioned whether direct competition alone was sufficient grounds for awarding an accounting. <u>Id.</u> at 317. The <u>Eckel</u> court was concerned that as opposed to infringement cases, the "assumption that defendant's profits correspond to the plaintiff's lost sales may be unfounded in a false advertising or unfair competition case." <u>Id.</u> "Without more," it continued, "the court cannot assume that the defendant's sales would have gone to the plaintiff but for the false advertisement or deceptive act." <u>Id.</u> To prevent a windfall to the plaintiff, the <u>Eckel</u> court proposed limiting the remedy of an accounting only to those cases "in which there was a false comparative advertisement or deceptive act that would lead consumer's [sic] to mistake the defendant's product for the plaintiff's." <u>Id.</u> at 318.

Nestle seizes on this language to support a per se rule prohibiting an accounting in all non-comparative advertising cases. I do not read Eckel so expansively. The Eckel court rightly highlighted a problem of causation that can arise in false advertising claims. A plaintiff and defendant may be in direct competition, but the defendant's alleged false advertising may not have caused harm directly to the plaintiff. Id. at 318.

Eckel's suggestion of limiting damage awards to comparative false advertising and palming off cases prevents providing the

plaintiff with a windfall in cases in which the plaintiff has proven direct competition, but no actual harm or causation.

Id. Where a plaintiff has alleged actual damages and causation, as is the case here, the concern for a windfall is significantly lessened. Thus, Eckel's bright-line rule is inapplicable.

Nestle's reliance on ALPO, 913 F.2d 958, is also misplaced. In ALPO, two dog food producers sued each other for false advertising under the Lanham Act. After a 61-day bench trial, the district court enjoined both parties from making the advertising claims found to be false and, pursuant to §1117(a), awarded ALPO over \$10 million, Ralston's estimated profits from its wrongdoing. Id. at 961. Then-Circuit Judge Clarence Thomas writing for the Court of Appeals, vacated the monetary damages on the basis that, in the absence of willful targeted wrongdoing, damages based upon profits was improper. Id. Judge Thomas relied on the D.C. Circuit's holding in Foxtrap, Inc. v. Foxtrap, Inc., 671 F.2d 636, 641 (D.C.Cir. 1982), which required a showing of bad faith as a prerequisite to an award based upon defendant's profits.

ALPO does not support Nestle's position for two reasons.

First, Foxtrap acknowledged that where no bad faith was shown, courts nevertheless may award profits where plaintiff has made a showing of actual competition, such as diversion of sales. Id.

 $^{^{16}}$ I note that the <u>Eckel</u> court denied monetary damages because the plaintiff failed to show actual harm and causation. <u>Eckel Indus., Inc. v. Primary Bank</u>, 26 F. Supp. 2d 313, 318 (D.N.H. 1998).

at 642 n9. Second, and more important, the First Circuit in Tamko, 282 F.3d at 36, explicitly rejected Foxtrap's requirement of bad faith in cases of direct competition. It bears repeating that "it has been [the First C]ircuit's rule that an accounting of defendant's profits where the products directly compete does not require fraud, bad faith, or palming off." Tamko, 282 F.3d at 36.

I also find inapplicable the other cases upon which Nestle Riggs Investment Management Corp. v. Columbia Partners, <u>L.L.C.</u>, 966 F. Supp. 1250, 1270 (D.C.Cir. 1997), relies on Foxtrap, 671 F.2d at 641, which as explained above, neither precludes an award of profits nor controls in this circuit. McNeilab, Inc. v. American Home Prods. Corp., 848 F.2d 34, 38 (2d Cir. 1988) holds that because the injury in non-comparative advertising cases accrues equally to all competitors, "some indication of actual injury and causation" is required before awarding damages under the Lanham Act -- a position consistent with the law of the First Circuit. Healthpoint, Ltd. v. Stratus Pharmaceuticals, Inc., 273 F.Supp.2d 871 (W.D.Tex. 2001) involved a question that is not at issue in this case: the applicability of a presumption of causation and injury in non-comparative advertising. Harper House, Inc. v. Thomas Nelson, Inc., 889 F.2d 197, 209 (9th Cir. 1989), was concerned with a similar presumption and is similarly inapposite.

Burndy Corp. v. Teledyne Indus., Inc., 748 F.2d 767, 772 (2d Cir. 1984), appears to support Nestle's argument in that it

states that the remedy of an accounting

appears to have been limited to situations in which the defendant's profits represent unjust enrichment derived from diversion of business that clearly would otherwise have gone to the plaintiff, such as in instances where the defendant palmed off its goods as made by the plaintiff or otherwise infringed the plaintiff's rights rather than engaged simply in false advertising of the defendant's own product.

However, properly read, this statement merely echos the views expressed in <u>Eckel</u>, 26 F. Supp. 2d at 317, and <u>McNeilab</u>, 848 F.2d at 38, that a false advertising plaintiff must show some actual harm and causation, such as "diversion of business that clearly would otherwise have gone to the plaintiff." <u>Burndy</u>, 748 F.2d at 772.

As Magistrate Judge Sorokin correctly stated, in order to survive a motion for judgment on the pleadings, Vermont Pure must allege facts which, if true, would entitle it to disgorgement of profits. McCord v. Horace Mann Ins. Co., 390 F.3d 138, 141 (1st Cir. 2004). Vermont Pure has alleged that "as a direct and proximate result of Nestle's false and misleading advertising concerning its Poland Spring bottled water," it suffered damages by the direct diversion of sales. Amended Complaint at ¶ 46. In support of this allegation, it further alleges that Nestle captured a significant market share through false advertising, and as a direct competitor, Vermont Pure suffered damages. Id. at ¶ 1. I agree with Magistrate Judge Sorokin that this is sufficient to survive a motion for judgment on the pleadings. Consequently, I find Nestle's motion to strike, or in the

alternative for judgment as a matter of law, as to Vermont Pure's claim for disgorgement of profits to be unfounded.

III. CONCLUSION

For the reasons set forth more fully above, I DENY Nestle's motion [98] to dismiss the quality claims and its motion [74] to strike Vermont Pure's claim for disgorgement of profits; and I ALLOW Vermont Pure's motion [67] for reconsideration of the "spring water" claims as to state causes of action. In connection with allowance to proceed on state "spring water"

claims, I direct Vermont Pure to file a Second Amended complaint on or before April 14, 2006 setting forth with particularity the precise state law "spring water" claims it intends to press.

/s/ Douglas P. Woodlock
DOUGLAS P. WOODLOCK
UNITED STATES DISTRICT JUDGE